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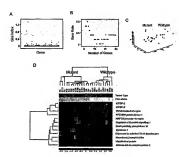
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[Continued on next page]

(54) Title: MOLECULAR SIGNATURE OF THE PTEN TUMOR SUPPRESSOR



(57) Abstract: The present invention relates to the identification a molecular signature for PTEN tumor suppressor. The molecular signature comprising a gene or genes that are of use for diagnosis, prognosis, drug research and development and therapeutics. Specifically, the present invention relates to identication of IGFBP2 gene, its mRNA and/or protein products that closely associate with PTEN mutations. The present invention further demonstrates that IGFBP2 expression is negatively regulated by PTEN and Akt activation, that IGFBP2 just a functional role in the PTEN signaling and is required for Akt transformation. The use of IGFBP2 gene, its gene product such as its RNA transcript, protein and molecular probes in diagnossis, reognosis, drug discovery and validation and therapeutic target and thisrapeutics is also contemplated.

2005/059109 A3 III

- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments
- For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.
- (88) Date of publication of the international search report: 28 September 2006

INTERNATIONAL SEARCH REPORT

International application No.

			PCT/US04/422		
A. CLAS					
USPC: According to	435/6 International Patent Classification (IPC) or to both na	ational	classification and IPC		
B. FIEL	DS SEARCHED				
Minimum documentation searched (classification system followed by classification symbols)					
U.S. : 42	35/6, 91.1, 91.2, 183; 436/94; 536/23.1, 24.3, 24.33, 2	5.3	,		
Documentati	on searched other than minimum documentation to the	exten	t that such documents are included	in the fields scarched	
	ata base consulted during the international search (namontinuation Sheet	e of da	ata base and, where practicable, sea	rch terms used)	
C. DOC	UMENTS CONSIDERED TO BE RELEVANT		-		
Category *	Citation of document, with indication, where a US 2006/0127902 A1 (MADDEN et al) 15 June 200			Relevant to claim No.	
	document, especially see [0008] in pages 4-6 and cl	aim 22	in pages 139-161		
	documents are listed in the continuation of Box C.		See patent family annex.		
"A" document	Special categories of cited documents: 'document defining the general state of the art which is not considered to be of particular relevance		later document published after the ini- date and not in conflict with the appli principle or theory underlying the inv	cation but cited to understand the ention	
E" carlier app	plication or patent published on or after the international filing date	"X"	document of particular relevance; the considered novel or cannot be consid when the document is taken alone		
	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as pecified)		document of particular relevance; the considered to involve an inventive ste with one or more other such documen	D when the document is combined	
	referring to an oral disclosure, use, exhibition or other means		obvious to a person skilled in the art	out outstand being	
P" document published prior to the international filing date but later than the priority date claimed		"&" 	document member of the same patent	family	
Date of the actual completion of the international search		Date	of mailing of the international sea	ch report	
26 June 2006 (26.06.2006)			O / A()(5 2006		
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	(571) 273-3201		V		

Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Form PCT/ISA/210 (second sheet) (April 2005)

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INTERNATIONAL SEARCH REPORT

International application No.

	101/030442236
Box No. II	Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)
This internati	onal search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1.	Claims Nos: because they relate to subject matter not required to be searched by this Authority, namely:
2.	Claims Nos: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3.	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box No. III	Observations where unity of invention is lacking (Continuation of item 3 of first sheet)
This Internation	onal Searching Authority found multiple inventions in this international application, as follows: ntinuation Sheet
3.	As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims. As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of any additional fees. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos:
	payment of a protest fee. The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
	No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet(2)) (April 2005)

INTERNATIONAL SEARCH REPORT

International application No. PCT/US04/42258

BOX III. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claims 1-20, drawn to a method of profiling a tumor/cancer in human tissue specimens wherein said gene is insulin-like growth factor binding protein 2.

Group II, claims 21-43, drawn to a method of screening a compound inhibits cancer cell growth.

Group III, claims 44-58, drawn to an assay kit of profiling a tumor/cancer in human tissue specimens.

Group IV, claims 59 and 60, drawn to a therapeutic useful antibody against insulin-like growth factor binding protein 2 or IGERP2

Note that, since claims 8 and 9 has a lot of genes which have different structures and different functions, Group I is divided to many different subgroups each subgroup has different spens, applicant requires to select a single gene for the examination. Applicant is advised that examination will be restricted to only elected gene and should not to be construed as a species election.

The inventions listed as Groups I to IV do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Groups I and II do not relate to a single general inventive concept under PCT Rule 13.1 because they lack the same or corresponding special technical features. For example, profiling a tumor/cancer in human tissue specimens in claim 1 of Croup I is not required for Group II white screening a compound that inhibits cancer cell growth in claim 21 of Group II is not required.

Groups I and III do not relate to a single general inventive concept under PCT Rule 13.1 because the technical feature linking
Groups I and III is not special. For example, an assay kit of Group III is not a contribution over the prior art since a kit containing
antibodies assainst HC ex-39 in US Patent No. 5.75.66 (published on March 10, 1998) reads an assay kit of Group III.

Groups I and IV do not relate to a single general inventive concept under PCT Rule 13.1 because the technical feature linking Groups I and III is not special. For example, profiling a tumor/cancer in human tissue specimens in claim 1 of Group I is not required for Group IV while a therapeutic useful antibody against insulin-like growth factor binding protein 2 in claim 59 of Group IV is not required for Group IV.

Groups II and III do not relate to a single general inventive concept under PCT Rule 13.1 because the technical feature linking Groups II and III is not special. For example, an assay kit of Group III is not a contribution over the prior art since a kit containing antibodies against HC gp.39 in US Patent No. 5.72.06 (published on March 10, 1998) reads an assay kit of Group III is

Groups II and IV do not relate to a single general inventive concept under PCT Rule 13.1 because they lack the same or corresponding special technical features. For example, extreming a compound that inhibits cancer cell growth in claim 21 of Group II is not required for Group IV while a therapeutic useful antibody against insulin-like growth factor binding protein 2 in claim 59 of Group IV is not required for Group II.

Groups III and IV do not relate to a single general inventive concept under PCT Rule 13.1 because they lack the same or corresponding special technical features. For example, an assay kit in claim 44 of Group III is not required fromp IV while at therapeutic useful antibody against insulin-like growth factor binding protein 2 in claim 59 of Group IV is not required for Group III.

	International application No.					
INTERNATIONAL SEARCH REPORT						
INTERNATIONAL SEARCH REPORT	PCT/US04/42258					
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Continuation of B. FIELDS SEARCHED Item 3:						
STN and EAST						
Scarch terms: cancer or tumor, express\$, gene, human, compound or reagent or substance, and insulin-like growth factor binding protein						